

**PAGE NUMBERING SEQUENCE IN THIS SECTION IS
INCONSISTENT**

INITIALS BAC.
DATE 7/28/05

APPENDIX A
REFERENCES

APPENDIX A

REFERENCES

1. U. S. Atomic Energy Commission, AEC Manual, Chapter 0510-01, "Prevention, Control, and Abatement of Air and Water Pollution by Federal Activities."
2. Idem, AEC Manual, Chapter 0511-01, "Radioactive Waste Management."
3. Idem, Plan for the Management of AEC-Generated Radioactive Wastes, Wash-1202 (January 1972).
4. Idem, Idaho Operations Office, NRTS Waste Management Plan for FY-1974, IDO-10051(4) (December 1973).
5. U. S. Atomic Energy Commission, "Prevention, Control, and Abatement of Air and Water Pollution at Federal Facilities," Federal Register, 35, 25 (February 5, 1970).
6. Letter, F. P. Baranowski, Director, Division of Production, to Managers, Richland, Idaho, and Savannah River Operations Offices, "Waste Management Environmental Impact Statements" (August 14, 1973).
7. U. S. Atomic Energy Commission, AEC Manual, Chapter 0524, "Standards for Radiation Protection."
8. Idem, Idaho Operations Office, Radioactive Waste Management Information 1974 Summary and Record-to-Date, IDO-10054 (74).
9. Argonne National Laboratory - West, Facility Waste Description Argonne West (December 1973) [Draft].
10. Idem, EBR-II System Design Descriptions, Volume I, General Facilities -- Argonne National Laboratory (June 1972).
11. L. J. Koch et al, Hazards Summary Report, Experimental Breeder Reactor II (EBR-II), ANL-5719 and Addendum (May 1957).
12. G. A. Freund et al, Design Summary Report on the Transient Reactor Test Facility, ANL-6034 (June 1960).
13. Argonne National Laboratory - West, Reactor Analysis and Safety Division and TREAT Operations, TREAT Baseline Description Document, ANL/RAS 72-73 (Draft).
14. H. Lawroski et al, Final Safety Analysis Report on the Zero Power Plutonium Reactor (ZPPR) Facility, ANL-7471 (June 1972).
15. Argonne National Laboratory - West, ZPPR Facility Operating and Maintenance Manual.

16. Idem, Fuel Manufacturing Operations Manual (unpublished).
17. Idem, HFEF Feasibility and Cost Study, Volume I, Conceptual Design Description (February 1968).
18. Idem, HFEF Title I Design Report and Cost Estimate (April 1969).
19. J. C. Hesson et al, Description and Proposed Operation of the Fuel Cycle Facility for the Second Experimental Breeder Reactor (EBR-II), Argonne National Laboratory - West (April 1963).
20. R. R. Smith et al, Exposed Fuel Calibration Study in EBR-II, Second Series, ANL-7558 (January 1970).
21. R. R. Smith and C. B. Doe, Published Failure Simulation Tests in EBR-II, ANL-7067 (December 1966).
22. Idem, Fission Product Monitoring in EBR-I, Mark IV, ANL-6788 (January 1964).
23. K. B. Purges, Reactor Physics Division Report, EBR-II Fuel Element Failure Detectors, ANL-7010 (January 1965).
24. G. S. Brunson, "On-Line Noble Gas Fission Product Monitoring in Experimental Breeder Reactor II," Nuclear Technology, 10 (January 1971) p 33.
25. K. G. Purges and A. DeVolpi, "Design Improvements and Sensitivity Tests of the FERD System for EBR-II," Reactor Physics Division Annual Report, ANL-7110 (December 1965).
26. Argonne National Laboratory - West, Drawing SW-1, "HFEF North and South Suspect and Liquid Waste" (October 1971).
27. B. J. Grady, Jr., Argonne National Laboratory - West, Site Engineering Section, Technical Support and Engineering Department, ANL-West Suspect Liquid Waste Baseline Description, Document No. W7500-1801-SA-00 (November 1972).
28. Argonne National Laboratory - West, Master Site Plan.
29. Idaho Board of Environmental and Community Services, Proposed Rules and Regulations for the Establishment of Standards of Water Quality and for Waste Water Treatment Requirements for Waters in the State of Idaho, (March 1973) [Revised].
30. H. V. Chamberlain, Safety Analysis Modifications of ICPP Dissolver and Vessel Off-Gas Systems, ICP-1011 (October 1973).
31. C. L. Bendixsen and G. F. Offutt, Rare Gas Recovery Facility at the Idaho Chemical Processing Plant, IN-1221 (April 1969).

32. L. T. Lakey and J. R. Bower (eds.), ICPP Waste Calcining Facility Safety Analysis Report, IDO-14620 (December 1963).
33. R. E. Girton et al, The Stack Monitor System at the Idaho Chemical Processing Plant, ICP-1034 (September 1973).
34. H. S. Cole, Disposal of Radioactive Liquid Organic Wastes at the ICPP, ICP-1007 (March 1972).
35. J. A. Wielang et al, The Fourth Processing Campaign in the Waste Calcining Facility, ICP-1004 (1972).
36. J. A. Wielang and W. A. Freeby, The Fifth Processing Campaign in the Waste Calcining Facility, ICP-1021 (1973).
37. Environmental Statement for the NWCF, Wash 1531 January 1974.
38. H. V. Chamberlain, Final Safety Analysis for ICPP Service Waste Diversion System, ICP-1020 (November 1973).
39. G. E. Lohse, Safety Analysis Report for the ICPP High-Level Solid Radioactive Waste Storage Facilities, ICP-1005 (January 1972).
40. U. S. Atomic Energy Commission, Environmental Statement, Loss-of-Fluid Test Facility, National Reactor Testing Station, Idaho, WASH-1517 (January 1973).
41. Idaho Air Pollution Control Commission, Rules and Regulations for the Control of Air Pollution in Idaho (1970).
42. U. S. Environmental Protection Agency, "National Primary and Secondary Ambient Air Quality Standards," Federal Register, 36, (84), Part II (1971) pp 8186-8201.
43. Aerojet Nuclear Company, LOFT Integral Test System Final Safety Analysis Report, Section 13.0 (to be published).
44. U. S. Atomic Energy Commission, Environmental Statement, Power Burst Facility, National Reactor Testing Station, Idaho, WASH-1514 (April 1972).
45. G. W. Hogg et al, A Survey of NRTS Waste Management Practices, ACI-104 (September 1971).
46. D. C. Hampson, "Radioactive Waste Management EBR-II Site, Argonne National Laboratory," prepared for presentation to the National Academy of Science Committee on Radioactive Waste Management, Idaho Falls, Idaho (April 17-18, 1969).
47. J. F. Kunze and P. L. Chase, Facility Operating Manual, Low Power Test Facility, CI-1188, Appendix A (December 1970).

48. J. E. Grund and B. E. Norton, SPERT I Low Enrichment Oxide Core Destructive Test Program Safety Analysis Report, IDO-16906 (August 1963).
49. C. R. Montgomery, SPERT II Reactor Facility, IDO-16888 (August 1963).
50. J. Dugone (ed.), SPERT III Reactor Facility, E-Core Revision, IDO-17036 (November 1965).
51. R. E. Heffner et al, SPERT IV Facility, IDO-16745 (February 1962).
52. H. L. Sletten, Organic Moderated Reactor Experiment Safeguards Summary, NAA-SR-2323 (February 1958).
53. Phillips Petroleum Company, Atomic Energy Division, Experimental Organic Cooled Reactor Safety Analysis Report, IDO-16820 (November 30, 1962).
54. R. O. Haroldsen et al, Safety Analysis Report, EBR-I, Mark IV, ANL-6411 (February 1963).
55. Argonne National Laboratory - West, BORAX III Operating Instructions.
56. Idem, BORAX IV Operating Instructions (April 1964).
57. G. R. Yansky et al, Climatology of the National Reactor Testing Station, IDO-12048 (1966).
58. G. A. Demarrais and N. L. Islitzer, The Diffusion Climatology of the National Reactor Testing Station, IDO-12051 (1959).
59. G. A. Demarrais, The Engineering Climatology of the National Reactor Testing Station, IDO-12004 (1958).
60. Idem, The Climatology of the National Reactor Testing Station, IDO-12003 (1958).
61. E. Walker, Subsurface Geology of National Reactor Testing Station, Idaho, USGS Bulletin 1133-E (1964).
62. R. L. Nace, Geography, Geology, and Water Resources of the National Reactor Testing Station, Idaho, Part 1, IDO-22033 (1956).
- 62a. R. L. Armstrong et al, "K-Ar Dating of Neogene Volcanic Rocks of the Snake River Plain, Idaho," American Journal of Science, 275, (March 1975) pp 225-251.
63. D. B. Hawkins and A. L. Short, Equations for the Sorption of Cesium and Strontium on Soil and Clinoptilolite, IDO-12046 (1965).
64. International Conference of Building Officials, Uniform Building Code, Volume I, 11th ed., Revised, Los Angeles, California (1961).

65. Idem, Uniform Building Code, 1970 ed., Volume I.
66. R. A. Eppley, Earthquake History of the United States, Part I, U. S. Department of Commerce (1965).
67. S. W. Nile, "The Hebgen Lake Earthquakes," 11th Annual Field Conference of the Billings Geological Society (1960) p 24.
68. E. Malde et al, Geological Investigation of Faulting Near the National Reactor Testing Station, Idaho, USGS open file report (1971).
69. A. H. Dahl and M. R. Niccum, "Preliminary Results of a Microseism Study for the Region Around the Snake River Plain," 12th Annual Engineering Geology and Soils Engineering Symposium, Idaho State University, Pocatello, Idaho (April 4-6, 1973).
70. J. T. Barraclough et al, Hydrology of the National Reactor Testing Station, Idaho, IDO-22049-USGS (1967).
71. P. H. Carrigan, Probability of Exceeding Capacity of Flood Control System at the National Reactor Testing Station, Idaho, USGS open file report (1971) 123 p.
72. R. D. Lamke, Stage Discharge Relations on Big Lost River within National Reactor Testing Station, Idaho, USGS open file report, IDO-22050, issued by U.S. Atomic Energy Commission, Idaho Falls, Idaho (1969) 29 p.
73. M. J. Mundorff, E. G. Crosthwaite, C. Kilburn, Ground Water for Irrigation in the Snake River Basin in Idaho, USGS Water Supply Paper 1654 (1964) 224 p., 6 pls., 54 figs., 14 tables.
74. R. F. Norvitch, C. A. Thomas, R. J. Madison, "Artificial Recharge to the Snake Plain Aquifer; an Evaluation of Potential and Effect," Water Information Bulletin No. 12, issued by Idaho Department of Reclamation, Boise, Idaho (1969) p 20.
75. R. L. Nace, J. W. Stewart, W. C. Walton et al, Geography, Geology, and Water Resources of the National Reactor Testing Station, Idaho -- Part 3, Hydrology and Water Resources, USGS open file report (1959) 253 p.
- 75a. J. T. Barraclough and R. G. Jensen, Hydrologic Data for the INEL-Site, Idaho, IDO-22055 (1926).
76. J. B. Robertson, R. Schoen, J. T. Barraclough, The Influence of Liquid Waste Disposal on the Geochemistry of Water at the National Reactor Testing Station, Idaho: 1952-70, USGS IDO-22053, issued by U.S. Atomic Energy Commission, Idaho Falls, Idaho (1974) 231 p.
- 76a. E. H. Walker, Subsurface Geology of the National Reactor Testing Station, Idaho, USGS Bulletin 1133-E (1964).

- 76b. R. L. Nace et al, 1975 Generalized Geologic Framework of the National Reactor Testing Station, Idaho, USGS professional paper 725-B (1975).
77. R. O. Harris and N. E. West, "Vegetation Patterns of the National Reactor Testing Station, Southeastern Idaho, Northwest Science 47, 30-43.
78. P. H. Wilks et al, Comprehensive Technical Report - ANP, APEX-921 (1972).
79. J. R. Horan (ed.), AEC Health and Safety Division Annual Report - 1958, IDO-12012 (1959).
80. U. S. Atomic Energy Commission, Idaho Operations Office, AEC Health and Safety Division Annual Report - 1959, IDO-12014 (October 1960).
81. Idem, AEC Health and Safety Division Annual Report - 1960, IDO-12019 (September 1961).
82. Idem, AEC Health and Safety Division Annual Report - 1961, IDO-12021 (June 1961).
83. Idem, AEC Health and Safety Division Annual Progress Report, IDO-12033 (September 1963).
84. R. E. Baker, Second Limited Melt Experiment, Final Report, APEX-714 (1961).
85. General Electric Company, Atomic Products Division, Aircraft Nuclear Propulsion Department, Summary Report of HTRE #3 Nuclear Excursion, APEX-509 (1959).
86. W. L. Ginkel et al, Nuclear Incident at the Idaho Chemical Processing Plant, IDO-10035 (1960).
87. J. J. DiNumno et al, Calculation of Distance Factors for Power and Test Reactor Sites, TID-14844 (1962).
88. Joint Committee on Atomic Energy, Congress of the United States, SL-1 Accident - Atomic Energy Commission Investigation Board Report, (June 1961).
89. Capt. A. Nelson Tardiff, Some Aspects of the WTR and SL-1 Accidents, IDO-19308 (April 1962).
90. U. S. Atomic Energy Commission, Idaho Operations Office SL-1 Report Task Force, Nuclear Incident at the SL-1 Reactor, IDO-19302 (January 1962).
91. General Electric Company, Final Report of SL-1 Recovery Operation, IDO-19311 (July 27, 1962).

92. R. W. Miller et al, Report of the SPERT I Destructive Test Program on an Aluminum, Plate-Type, Water-Moderated Reactor, IDO-16883 (June 1964).
93. D. F. Bunch, SPERT I Destructive Test Series Environmental Monitoring and Research Studies, IDO-12039 (January 1965).
94. O. L. Cordes et al, Radiological Aspects of SNAPTRAN 2/10A-3 Destructive Test, IDO-17038 (January 1965).
95. O. L. Cordes et al, Radiological Aspects of the SNAPTRAN-2 Destructive Tests, IDO-17203 (February 1967).
96. C. A. Hawley, Jr. (ed.), Controlled Environmental Radioiodine Tests at the National Reactor Testing Station -- 1965 Progress Report, IDO-12047 (February 1966).
97. D. F. Bunch (ed.), Controlled Environmental Radioiodine Tests Progress Report Number Two, IDO-12053 (August 1966).
98. Idem, Controlled Environmental Radioiodine Tests -- Progress Report Number Three, IDO-12063 (1967).
99. J. D. Zimbrick and P. G. Volleque' (eds.), 1967 CERT Progress Report Controlled Environmental Radioiodine Tests at the National Reactor Testing Station -- Progress Report Number Four, IDO-12065 (December 1968).
100. E. P. Hardy, Jr., Health and Safety Laboratory Environmental Quarterly, USERDA Report HASL-291, 1975.
101. Milton W. Meyer et al, Strontium-90 on the Earths Surface IV, TID-24341 (May 1968).
102. U. S. Atomic Energy Commission, Idaho Operations Office Health Services Laboratory, National Reactor Testing Station, Idaho (unpublished data).
103. J. H. Osloond et al, Radioactive Waste Disposal Data for National Reactor Testing Station and Supplements, IDO-12040 (April 1965).
104. B. L. Schmalz, Radionuclide Distribution in Soil Mantle of the Lithosphere as a Consequence of Waste Disposal at the National Reactor Testing Station, IDO-10049 (October 1972).
- 104a. J. B. Robertson, Numerical Modeling of Subsurface Radioactive Solute Transport from Waste Seepage Ponds at the INEL, USGS, 76-717, IDO-22057 (1976).
- 104b. W. L. Polzer et al, Special Analyses for Plutonium and Americium in Water from the Snake River Plain Aquifer, IDO-12081 (November 1976).

105. J. B. Robertson and J. T. Barraclough, "Radioactive and Chemical Waste Transport in Groundwater at the NRTS -- 20 Year Case History and Digital Model," a paper for presentation to International Symposium on Underground Waste Management and Artificial Recharge, New Orleans, Louisiana September 26, 1973.
106. J. Echo and D. Hawkins, "Algal Influence on Radionuclides in Settling Ponds," Nature, 209 (March 1966) p. 115.
107. U. S. Atomic Energy Commission, Idaho Operations Office, 1963 Annual Report, Health and Safety Division, IDO-12037 (August 1964).
108. J. C. Commander, Nonradioactive Waste Oil Disposal Study, CI-1212 (February 1971).
109. U. S. Public Health Services, Drinking Water Standards (1962).
110. J. T. Barraclough et al, Hydrology of the Solid Waste Burial Ground as Related to the Potential Migration of Radionuclides, Idaho National Engineering Laboratory, IDO-22056 (August 1976).
- 110a. W. H. Burgus and S. E. Maestas, The 1975 RWMC Core Drilling Program, IDO-10065 (July 1976).
- 110b. J. L. Harness et al, Onsite Environmental Surveillance Report for the INEL Radioactive Waste Management Complex, Annual Report 1974, TREE-1014 (December 1976).
111. NAS-NRC Committee, The Effects on Populations of Exposure to Low Levels of Ionizing Radiation, The BEIR Report, Washington, D.C.: National Academy of Sciences (1972).
112. United Nations Scientific Committee on the Effects of Atomic Radiation, Ionizing Radiation: Levels and Effects, New York: United Nations (1972).
113. Environmental Protection Agency, Environmental Radiation Dose. Commitment: An Application to the Nuclear Power Industry, EPA-520/4-73-002, Washington, D.C.: U. S. Government Printing Office (1974).
114. Doris Karlsson (ed.), Safety Review Document for the RWMC (to be published).
115. Idaho Water Resources Board, Interim State Water Plan - Preliminary Report - 1972.
116. M. J. Mundroff et al, Ground Water for Irrigation in the Snake River Basin in Idaho, USGS Water Supply Paper (1964).
117. Idaho Code - Title 42, Chapter 39.

118. Idaho Department of Environmental and Community Services (E&CS), "Water Quality Standards. . . Requirements," adopted by resolution of Board of E&CS (June 28, 1973).
119. "The Shallow Land Burial of Low-Level Radioactively Contaminated Solid Waste," Library of Congress Catalog Card Number 76-56928. Available from Printing and Publishing Office, National Academy of Sciences, 2101 Constitution Avenue, N.W., Washington, D.C. 20418.

APPENDIX B
GLOSSARY OF TERMS
AND ABBREVIATIONS

APPENDIX B

GLOSSARY OF TERMS

AND ABBREVIATIONS

The explanation of terms used in the text is prepared according to the general usages developed during operations on the INEL site. Where possible, the explanations are developed also in accordance with established standards.

Acronyms, a word identified by first letters of a series of words, are also listed.

A

acre	a unit of area measurement equal to 43,560 ft ²
activation	the induction of radioactivity in material by irradiation with neutron radioactive material, a radiation generating machine, or a nuclear reactor
activity	a measure of the rate at which a material is emitting nuclear radiations, usually given in terms of the number of nuclear disintegrations occurring in a given quantity of material over a unit of time; the standard unit of activity is the curie (Ci)
alpha radiation	an emission of particles (helium nuclei) from a material undergoing nuclear transformation; the particles have a nuclear mass number of four and charge of plus two
alpha waste	waste material which is contaminated by radionuclides which emit alpha particles, particularly transuranic elements
aquifer	a subsurface formation containing sufficient saturated permeable material to yield significant quantities of water
aseismic	not subject to earthquakes
atomic number	the number of protons in the nucleus of each chemical element
alluvial fan	rock deposit laid down by streams flowing from mountains into lowland regions

B

background radiation	the level of radioactivity in an area, which is produced by sources other than the one of specific interest; in the INEL region, the background radiation is produced by naturally occurring radioactive materials in the crust of the earth, cosmic radiations, and the fallout from nuclear weapons tests
beta radiation	essentially weightless charged particles (electrons and positrons) emitted from the nucleus of atoms undergoing nuclear transformation
bioconcentration (bioaccumulation)	the process whereby an organic system selectively removes an element from its environment and accumulates that element in a higher concentration
biological oxygen demand (BOD)	a measure of the organic pollution of water, determined by the extent to which bacteria and other contained organisms in a water sample will use dissolved oxygen in a given period of time; therefore, a measure of the residual oxygen in the water for use by other organisms such as fish
biosphere	living beings together with their environment
biota	the plant and animal life of a region
biotic	caused by living organisms
blowdown	cooling tower effluent water with a higher dissolved solids content than natural water
body burden	the amount of specified radioactive material or the summation of the amounts of various radioactive materials present in an animal or human body at the time of interest
burial ground	an area specifically designated for the subsurface disposal of solid waste or excess materials

C

calcination	solidification method for disposal of liquid wastes involving (a) atomizing and coating of liquid on small granular solids and (b) heating to drive off moisture
cask	a container designed for the transporting of radioactive materials, the design usually includes special shielding, handling, and sealing features to provide positive containment of the materials and to minimize exposure of personnel
chemical oxygen demand (COD)	a measure of the extent to which all chemicals contained in a water sample use dissolved oxygen in a given period of time; therefore, a measure of residual dissolved oxygen in the water available for use by organisms such as fish
coliform (count, number)	a measure of the bacterial content of water; a high coliform count indicates potential contamination of a water supply by human waste
concentration guide	the average concentration of a radionuclide in air or water to which a worker or member of the general population may be continuously exposed without exceeding appropriate radiation dose standards (see maximum permissible concentration)
contamination (contaminated material)	the deposition, solvation, or infiltration of radionuclides on or into an object, material, or area whereupon the area, material, or object is considered "contaminated"; loose contamination is considered removable by conventional decontamination methods; fixed contamination is not removable by conventional decontamination methods
controlled area	any specific region of the INEL into which entry by personnel is regulated by physical barrier or procedure
counts per minute (cpm)	the number of events per unit time recorded by an instrument designed to detect radioactive particles; especially used to indicate the relative amount of radioactive contamination

critical	the condition in which a material is undergoing nuclear fission at a self-sustaining rate; the critical mass of a material is that amount which will self-sustain nuclear fission when placed in an optimum arrangement in its present form; the minimum critical mass is the amount of a fissile isotope that will self-sustain nuclear fission when placed in optimum conditions
criticality safety	those procedures and understandings necessary to the handling of fissile materials in a manner that will prevent them from reaching a critical condition
curie (Ci)	a unit of radioactivity defined as the amount of a radioactive material that has an activity of 3.7×10^{10} disintegrations per second (d/s); millicurie (mCi) = 10^{-3} curie; microcurie (μ Ci) = 10^{-6} curie; nanocurie (nCi) = 10^{-9} curie; picocurie (pCi) = 10^{-12} curie; femtocurie (fCi) = 10^{-15} curie
D	
daughter products	the nuclides formed by the radioactive disintegration of a first nuclide (parent)
deactivated	the condition of a facility or disposal site where steps have been taken to preclude further operation or the further addition of waste materials
decay chain	the sequence of radioactive disintegrations in succession from one nuclide to another until a stable daughter is reached
decay heat	the thermal energy produced in a material by its own radioactive disintegrations
decontamination	the selective removal of radioactive material from a surface or from within another material
decommissioning	the process of removing a facility or area from operation and decontaminating and/or disposing of it or placing it in a condition of standby with appropriate controls and safeguards

disintegrations per minute (dpm)	the number of radioactive decay events occurring per unit time in a given amount of material
disposal	the planned release or placement of waste in a manner that precludes recovery or its placement in a manner which is considered permanent so that recovery is not provided
dose	a general term indicating the amount of energy absorbed from incident radiation by a specified mass
dose commitment	the integrated dose which results from an intake of radioactive material when the dose is evaluated from the beginning of intake to a later time (usually 50 years); also used for the long-term integrated dose to which people are considered committed because radioactive material has been released to the environment
drum	a metal or composition cylindrical container used for the transportation, storage, and disposal of waste materials
E	
ecology	that branch of biological science which deals with the relationships between organisms and their environment
ecosystem	an assemblage of biota (community) and habitat
environmental surveillance	a program to monitor the impact on the surrounding region of the discharges from industrial operations
evapotranspiration	the combined loss of water from soil by evaporation from the soil and from the surfaces of plant structures
excursion	a sudden rapid increase of power produced when a reactor or other system of fissile material undergoes a sudden increase in reactivity
exposure	the condition of being made subject to the action of radiation
extraction	a chemical process for selectively removing materials from solutions

F

fallout	those radioactive materials deposited on the earth's surface and in the atmosphere following the detonation of nuclear weapons
fast flux (fast neutron)	a stream of neutrons having energies (velocities) near that imparted to them by a fission event; when applied to nuclear reactors, refers to those using high velocity neutrons to cause successive fission events
fiissile	material capable of undergoing fission by any process
fissionable	material capable of undergoing fission by slow neutrons
fission (nuclear)	the division of a nucleus into two nuclides of lower mass, usually accompanied by the expulsion of gamma rays and neutrons
fission products	the nuclides formed by the division of a heavier nucleus; most usually in a nuclear reactor
foodchain	a linear sequence of successive utilizations of nutrient energy by a series of species
foodweb	the concept of nutrient energy transfer (including decomposition) between species in an ecosystem
fuel (nuclear, reactor)	fissionable material used as the source of power when placed in a critical arrangement in a nuclear reactor
fuel separation (fuel reprocessing)	processing of irradiated (spent) nuclear reactor fuel to recover useful materials as separate products, usually separation into plutonium, uranium, and fission products

G

gamma radiation	electromagnetic energy emitted in the process of a nuclear transition
gamma scan	process of measuring the energy spectrum of the gamma rays emitted by a material in order to determine its constituent nuclides

gastrointestinal dose (GI)	the dose to the stomach and lower tract of humans and animals via external exposure or via internal transport of radioactive material
GeigerMuller Tube	a gas-filled tube used to detect radiation events by the ionization pulse produced in the gas; used on a GM counter
groundwater	water which exists or flows below the surface (within the zone of saturation)
H	
habitat	the abiotic characteristics of the place biota live
half-life	the time required for the activity of a radionuclide to decay to half its value; used as a measure of the persistence of radioactive materials, each radionuclide has a characteristic constant half-life
high efficiency particulate air filter (HEPA)	an air filter capable of removing at least 99.97% of the particulate material in an air stream
high-heat waste	liquid radioactive waste which generates sufficient fission product decay heat to cause self-boiling and self-concentration
high-level liquid waste	the aqueous wastes resulting from the operation of the first-cycle extraction system, or equivalent concentrated wastes from subsequent extraction cycles; or equivalent wastes from a process not using solvent extraction, in a facility for processing irradiated reactor fuels
hood	a canopy and exhaust duct used to confine hazardous materials in order to reduce the exposure of industrial workers
hypothetical maximum individual (max man)	a postulated person who is assumed to receive the maximum credible radiological dose through each of the exposure pathways from the source being considered

I

inactive	the condition of a facility or disposal site which is not presently being operated or to which materials are not being added
inversion	a condition where temperature increases with height in the atmosphere
ion exchange	process for selectively removing a constituent from a waste stream by reversibly transferring ions between an insoluble solid and the waste stream; the exchange medium (usually a column of resin or soil) can then be washed to collect the waste or taken directly to disposal; for example, a hot water softener
irradiation	exposure to radiation by being placed near a radioactive source; usually in the case of fuel materials, being placed in an operating nuclear reactor
isopleth	a line connecting all points of equal air concentration in plotting that air concentration at ground level as a function of direction and distance from a point of release
isotope	nuclides with the same atomic number, (i.e., the same chemical element) but with different atomic masses; although chemical properties are the same, radioactive and nuclear properties may be quite different for each isotope of an element

L

leaching pond	an excavation used for the disposal of liquids so that the soil will remove contaminants while allowing water and other solvents to pass through
lithologic	pertaining to the characteristics and study of rocks
long-lived isotope	a radioactive nuclide which decays at such a slow rate that a quantity of it will exist for an extended period; usually radionuclides whose half-life is greater than 3 years

low-level liquid waste

liquid wastes that can be discharged to the environment with assurance that persons will not be exposed to concentrations in excess of those prescribed in ERDAM-0524

M

man-rem

used as a unit of population dose; the average dose per individual expressed in rems times the population affected

maximum permissible concentration (MPC)

the average concentration of a radio-nuclide in air or water to which a worker or member of the general population may be continuously exposed without exceeding an established standard of radiation dose limitation

N

neutron

a particle existing in or emitted from the atomic nucleus; it is electrically neutral and has a mass approximately equal to that of a stable hydrogen atom

neutron activation

the process of irradiating a material with neutrons so that the material itself is transformed into a radioactive nuclide

nitrogen oxides (NO_x)

a mixture of nitrogen-oxygen containing compounds formed as gaseous waste effluents during the combustion of most fossil fuels

nuclear fission

see fission

nuclear radiation

particles and electromagnetic energy given off by transformations occurring in the nucleus of an atom

nuclear reactor

a device constructed of fissionable material such that a chain of fission events can be maintained and controlled to meet a particular purpose

nuclide

a species of atom having a specific mass, atomic number, and nuclear energy state

nucleus

the positively charged center of an atom

P

penetrating radiation

forms of radiant energy which are capable of passing through significant thicknesses of solid material; these usually include gamma rays, X rays and neutrons

permissible dose	that dose of ionizing radiation which, in the light of present knowledge, carries negligible probability of causing severe somatic injury or genetic effect
pH	a measure of the relative acidity or alkalinity of solution; a neutral solution has a pH of 7, acids have pH's of 7 to 1, bases have pH's of 7 to 14
phytoplankton	microscopic plants that live drifting in a body of water
population dose (population exposure)	the summation of individual radiation doses received by all those exposed to the source or event being considered
R	
rad	a special unit of measure for the absorbed dose of radiation; one rad equals 100 ergs absorbed per gram of material
radiation (ionizing)	particles and electromagnetic energy emitted by nuclear transformations which are capable of producing ions when interacting with matter; gamma rays and alpha and beta particles are primary examples in INEL waste
radiation survey	evaluation of an area or object with instruments in order to detect, identify, and quantify radioactive materials and radiation fields present
radioactive (decay)	property of undergoing spontaneous nuclear transformation in which nuclear particles or electromagnetic energy are emitted
radioiodines	isotopes of iodine which are radioactive
radionuclide	a nuclide which is radioactive
radiotoxicity	the property of a material of being able to adversely affect biological organisms through the mechanism of nuclear radiation
radwaste	waste materials which are radioactive
raptor	bird of prey
reactivity	a measure of the capability of a system to maintain criticality; systems with high reactivity are capable of undergoing rapid excursions

of increasing power; systems with low reactivity will undergo slower excursions; systems with negative reactivity will not become critical

reactor

a nuclear reactor

regolith

rock "waste" or surface mantle of unconsolidated rock debris

release limit
(release guide)

a control number which regulates the concentration or amount of radioactive material released to the environment in an industrial situation; usually dose to persons in the environment derived from environmental behavior of the released material so that the dose is kept below a selected control value

rem

a unit of measure for the dose of ionizing radiation which gives the same biological effect as one roentgen of X rays; one rem approximately equals one rad for X, gamma, or beta radiation

reprocessing

chemical processing of irradiated nuclear reactor fuels to remove desired constituents

retention basin

an excavated and lined area used to hold contaminated fluids until radioactive decay reduces activities to levels permissible for release

retired facility

a facility which has been shut down with no intentions of restarting and which has had appropriate controls and safeguards placed on it

roentgen

a unit of measure of ionizing electromagnetic radiation (X and gamma); one roentgen corresponds to the release by ionization of 83.8 ergs of energy per gram of air

rupture

a breach of the metal cladding of a production reactor fuel element thereby releasing radioactive materials to reactor cooling streams

S

sanitary landfill

a burial operation for the purpose of disposing of sanitary wastes

sanitary sewage	human wastes and other nonradioactive material for disposal to preserve public health
scintillation (counter)	light flashes produced in crystalline material by ionizing radiation; measurement of the level of activity of the source
seepage pond	an artificial body of surface water formed by discharge from INEL process operations
seismicity	the tendency for the occurrence of earthquakes
shielding	bulkheads, walls, or other constructions used to absorb radiation in order to protect personnel or equipment
short-lived isotope	a radioactive nuclide which decays so rapidly that a given quantity is transformed into its daughter products within a short period (usually those with a half-life of days or less)
smear	a means of measuring loose surface contamination on an object by wiping it with paper, gauze, etc., and then measuring the disintegrations per minute on the wipe with an instrument
solid wastes (radioactive)	either solid radioactive material or solid objects which contain radioactive material or bear radioactive surface contamination
sorptive capacity	the measure of a material's ability to sorb specific constituents from liquid as it passes through the material
stability (atmospheric)	a description of the atmospheric forces on a parcel of air following vertical displacement in an atmosphere otherwise in hydrostatic equilibrium; if the forces tend to return the parcel to its original level, the atmosphere is stable; if the forces tend to move the parcel further in the direction of displacement, the atmosphere is unstable; if the air parcel tends to remain at its new level, the atmosphere has neutral stability
storage	retention of radioactive waste in some type of man-made device, such as a tank or vault, in a manner permitting retrieval
sulfur oxides (SO_2/SO_3)	compounds formed as waste effluents during the burning of some fossil fuels

T

tank	a large metal container located underground for storage of liquid wastes
tank farm	an installation of interconnected underground containers (tanks) for storage of high-level waste
tectonic	pertaining to, or designating the rock structures resulting from deformation of the earth's crust
tracer	a radionuclide(s) or chemical introduced in minute quantities to a system or process for the purpose of using radiation or chemical detection techniques to follow the behavior of the process or system
transmutation	the process whereby one nuclide changes (or is changed) into another; usually by the addition of nuclear particles
transuranium	nuclides having an atomic number greater than that of uranium (i.e., greater than 92)
transmissivity	a coefficient relating the volumetric flow through a unit width of groundwater to the driving force (hydraulic potential); a function of both the porous medium, fluid properties, and saturated thickness of the aquifer

V

vadose zone	the unsaturated region of soil between the ground surface and the water table
-------------	---

W

water table	upper boundary of an unconfined aquifer surface below which saturated groundwater occurs; defined by the levels at which water stands in wells that barely penetrate the aquifer
wind rose	a diagram designed to show the distribution of prevailing wind directions at a given location; some variations include wind speed groupings by direction

List of Abbreviations

ACC	Allied Chemical Corporation
AEC	Atomic Energy Commission
ANC	Aerojet Nuclear Company
ANL-W	Argonne National Laboratory-West
ANPP	Aircraft Nuclear Propulsion Program
AlW	Prototype Nuclear Surface Ship Propulsion System
APS	Atmospheric Protection System
ARA	Auxiliary Reactor Area
ARMF	Advanced Reactivity Measurement Facility
ATR	Advanced Test Reactor
ATRC	Advanced Test Reactor Critical Facility
BORAX	Boiling Water Reactor Experiment
cc	Cubic Centimetre
cm	Centimetre
CERT	Controlled Experimental Radioiodine Test
CFA	Central Facilities Area
CFR	Code of Federal Regulations
Ci	Curie
EBOR	Experimental Beryllium Oxide Reactor
EBR-I and -II	Experimental Breeder Reactors I and II
ECF	Expended Core Facility
EPA	Environmental Protection Agency
ERDA	Energy Research and Development Administration
ETR	Engineering Test Reactor
EFS	Experimental Field Station
EOCR	Experimental Organic Cooled Reactor

ETRC	Engineering Test Reactor Critical Facility
FAA	Federal Aviation Agency
g	gram (seismology, g. = gravity)
GCRE	Gas Cooled Reactor Experiment
HEPA	High Efficiency Particulate Air
HFEF	Hot Fuel Examination Facility
HSL	Health Services Laboratory
IET	Initial Engine Test
ICPP	Idaho Chemical Processing Plant
LMFBR	Liquid Metal Fast Breeder Reactor
LOCA	Loss-of-Coolant Accident
LPT	Low Power Test
LOFT	Loss-of-Fluid Test
μ Ci	Microcurie
mCi	Millicurie
mrad	Millirad
mREM	Millirem
mR/hr	Milli-Roentgen per hour
NOAA	National Oceanic and Atmospheric Administration
NRF	Naval Reactor Facilities
NRTS	National Reactor Testing Station
OMRE	Organic Moderated Reactor Experiment
PBF	Power Burst Facility
PEW	Process Equipment Waste
RaLa	Radioactive Lanthanum
RWMC	Radioactive Waste Management Complex
S5G	Prototype Nuclear Submarine Propulsion System

SDA	Subsurface Disposal Area
SLW	Prototype Nuclear Submarine Propulsion System
SL-1	Stationary Low Power Test Reactor
SNAP	Systems for Nuclear Auxiliary Power
SPERT	Special Power Excursion Reactor Test
TAN	Test Area North
TDA	Transuranic Disposal Area
TSA	Transuranic Storage Area
TRA	Test Reactor Area
TREAT	Transient Reactor Test Facility
TSF	Technical Support Facility
WEC	Westinghouse Electric Corporation
WCF	Waste Calciner Facility
WMIS	Waste Management Information System
ZPPR	Zero Power Plutonium Reactor

APPENDIX C
LISTING AND DISPOSITION OF PAST
INEL RELEASES

APPENDIX C

LISTING AND DISPOSITION OF PAST

INEL RELEASES

In addition to the major programmatic and accidental releases discussed in Section II.C, several other events of lesser importance have occurred at INEL. Brief discussions of those which were of significant environmental concern are presented chronologically in this appendix.

Many of the releases were related to programs no longer being conducted and/or presently decommissioned facilities. Others have resulted in upgrading of equipment, procedures, training, etc., to minimize the possibility of similar occurrences. Where the data were available, quantitative release values are given. In all the cases listed, no real or potential long-term commitment of resources or significant cumulative environmental impact exists.

①. BORAX-I Destruction

After a series of transient tests, the BORAX-I reactor was intentionally subjected to a rapid power excursion in July 1954. The excursion produced 135 MW of energy and resulted in melting most of the fuel plates and gross damage to the reactor. A description of the reactor is given in Section II.A.12.

Monitoring teams that were dispatched shortly after the excursion determined the trajectory of the released effluent to have been roughly in a southwesterly direction toward the INEL Radioactive Waste Management Complex. Surveys of the total fission product radioactivity of all the debris indicated that practically all of the fuel originally in the reactor could be accounted for within a radius of 350 ft around the original reactor location, with the majority in an area 350 x 200 ft just south of the reactor. Approximately 5,000 ft² of ground was covered with a 6-in. layer of gravel to reduce the dose rates to acceptable levels. During the past 20 years, the radiation levels in the area have decayed to near background levels.

2. ICPP Well Discharge

In July 1954 during a routine transfer to the discharge well, permissible levels were exceeded, and 51 μ Ci of long-lived fission product activity were released in 244,000 gallons of liquid.

③. EBR-I Fuel Meltdown

In November 1955 the EBR-I experienced an excursion which resulted in melting of some of the fuel elements. The reactor building was slightly contaminated from released fission product gases, but there were no significant personnel exposures or environmental releases.

4. Release to ICPP Discharge Well

In August 1956 approximately 1 Ci of long-lived fission product activity was released to the ICPP discharge well.

5. RaLa Release

In February 1958 activity from a RaLa process run was accidentally released into the building for approximately 5 min with minor exposures to personnel. An estimated 1 μ Ci of fission product activity, including iodine-131, was released to the environment.

6. BORAX-IV Fuel Defect Test

In March 1958 the BORAX-IV reactor was intentionally operated at a power of 2.4 MW with a large number of defective fuel elements. During the tests, radioactive fission products leaked from the fuel causing high radiation and building contamination. Environmental releases were primarily limited to cleanup operations.

7. Iodine Release at ICPP

In March 1958 approximately 1 Ci of iodine-131 was released to atmosphere during a routine waste transfer operation at ICPP.

8. Blower Failure at ICPP

In August 1958 an atmospheric release of approximately 10 μ Ci of long-lived fission product activity occurred at the ICPP owing to a blower failure.

9. Solvent Burner Release

In September 1958 approximately 0.25 Ci of long-lived particulate activity was released from the ICPP solvent burner via the main stack.

10. Fuel Element Cutting Facility Release

In October 1958 a filter failure resulted in a 1,200-Ci particulate release to atmosphere consisting primarily of long-lived fission products, from the fuel element cutting facility at ICPP which contaminated the land immediately adjacent to the facility.

11. Liquid Spill at ICPP

An accidental liquid spill in the fuel process building at ICPP required extensive decontamination activities from November 1958 through April 1959. As a result, a considerable quantity of liquid waste and decontamination solutions was processed in the plant waste system and approximately 100 Ci of long-lived fission product activity were packaged and sent to the INEL Radioactive Waste Management Complex.

12. Collection Tank Release

In December 1958 approximately 1 Ci of radioactive noble gases and iodine was released to the atmosphere from a liquid waste tank at ICPP owing to an leaking flange.

13. Liquid Release at ICPP

In December 1958 approximately 29 curies of activity (7 curies of strontium-90) was accidentally released to the ICPP disposal well from an unknown origin in the plant system.

14. ICPP Criticality

In October 1959 a criticality accident occurred in a 5,000-gallon tank when an air-sparging operation initiated an unanticipated siphoning of a uranium solution from a geometrically safe vessel into a geometrically unsafe 5,000-gallon tank containing some water. Roughly 350,000 curies of primarily short-lived gaseous fission products were released to the atmosphere via the plant stack (see Section II.C.11.c).

15. ICPP Criticality

In January 1961 a criticality occurred in the ICPP first-cycle product evaporator. A total of 5,200 Ci of primarily short-lived fission products were released to the environment.

16. Contamination of Tank Vault

In March 1962 at ICPP, high-level liquid waste siphoned from an underground waste tank into its secondary containment vault. No activity was released to the environment.

17. Tank Farm Incident

In May 1964 approximately 5 Ci of long lived fission product radioactivity were released when high-level liquid waste was accidentally discharged at ICPP during construction tie-in of new tanks.

18. Sodium Release and Fire in EBR-II Sodium Boiler Plant Building Control Room

In February 1968 approximately 80 gallons of sodium were accidentally released and immediately ignited at EBR-II. The sodium contained approximately 4 mCi of radioactive sodium-24. Approximately 0.4 μ Ci was subsequently released to the atmosphere.

19. TSF Waste Tank Overflow

In August 1969 a 10,000-gallon-capacity underground liquid catch tank at TSF overflowed through a vent pipe and contaminated the surrounding soil. The actual number of curies released was not determined; however, the contamination is localized to an approximate area of 8 ft², and current radiation levels are 5 R/hr at 1 ft.

20. ICPP Discharge Well

In September 1969 approximately 19 Ci of long-lived fission product activity were accidentally released to the ICPP discharge well.

21. ANL-W Area Drainage Ditch Contamination

In October 1969 radioactive liquid was accidentally discharged to an industrial waste drainage ditch at ANL-W. It was calculated that approximately 413 mCi were released consisting of 90% zirconium-niobium-95 and cerium-144, with traces of cobalt-60, antimony-124, and cesium-137. A survey in 1973 indicated the presence of approximately 9.4 mCi of activity remaining in the ditch.

22. ICPP Discharge Well

In December 1969 approximately 1.5 μ Ci of strontium-90 in excess of the discharge limits were inadvertently released to the ICPP discharge well.

23. Leak in TRA Liquid Waste Holding Tank

In July 1970 a leak occurred in a 9,000-gallon-capacity underground tank used to collect low- to intermediate-level liquid waste at TRA. Failure was due to corrosion, and the leak was detected by visual observation of a level indicator and presence of liquid in the tank sump. The liquid was transferred to another vessel, and the tank was removed to the INEL Radioactive Waste Management Complex. Approximately 0.578 Ci of radioactivity was discharged to the subsoil, consisting primarily of cerium-144, strontium-89, and strontium-90.

24. TSF Feed Tank Overflow

In February 1971 a 50,000-gallon-capacity underground liquid waste tank at TSF overflowed. Approximately 293 gallons, totaling about 0.88 Ci of long-lived fission products, were released to the surrounding soil.

25. TSF Evaporator Release

In September 1971 an accidental airborne release occurred from the TSF liquid waste evaporator. The release was determined to be approximately 266 μ Ci of cesium-137, 0.0142 μ Ci of strontium-90, and 0.0142 μ Ci of yttrium-90.

26. Ruthenium Release

In January 1972 approximately 1 Ci of particulate ruthenium-106 was released from the ICPP stack.

27. WCF Release

In May 1972 a leak from a decontamination line into a calcine transport air line resulted in a release to the ground of 10 Ci of long-lived fission product activity.

28. Leaking Underground Liquid Waste Lines

On April 4, 1974 approximately 20 to 30 yd³ of contaminated soil were discovered around a corroded carbon steel vent line from a liquid waste storage tank at the ICPP. The line was abandoned, capped, and a total of 275 yd³ of soil were removed to the INEL Radioactive Waste Management Complex (RWMC). The release totaled approximately 1,000 Ci of primarily Cs-137, Cs-134, Sr-90, and Eu-154.

On October 1, 1974 contaminated soil was discovered adjacent to an underground pipe which carried high-level liquid from the processing building to the high-level liquid waste storage tanks. The leak was caused by a 0.15.-in.-diameter hole inadvertently drilled through the pipe wall during installation of the encasement cover in 1955. Approximately 50 yd³ of soil containing an estimated 3,000 Ci of radioactivity had been relocated to the RWMC as of the end of October 1974. An estimated 3,000 Ci of radioactivity contained in an additional 50 yd³ of soil will be removed in the near future. This activity consists principally of Sr-Y-90, Ce-Pr-144, and Cs-137.

29. Subsurface Soil Contamination at ICPP

On September 18, 1975 an underground contaminated soil zone was discovered at the ICPP. The zone measured 150 by 20 ft along a backfilled area of a pipe trench at a depth generally between 12 and 25 ft. The source of contamination was a corroded carbon steel line in the radioactive liquid waste transfer system in the storage tank area. Approximately 14,000 gallons of radioactive liquid leaked from the corroded line. This represents an estimated 30,000 curies of radioactivity consisting primarily of Cs-137 and Sr-Y-90.

The corroded carbon steel line which was the source of the soil contamination was removed from service and capped off to prevent the possibility of future losses. The contaminated soil zone has been mapped and, if future excavation in the area becomes necessary, the excavated contaminated soil will be removed to the RWMC.

30. Diversion Valve Leak at ICPP

During a routine transfer of solution from a service waste diversion tank to an evaporator waste collection tank on January 16, 1976, solution was observed leaking from the flange on one of the diversion valves. Checks on the volume of waste received in the collection tank with that removed from the diversion tank indicate that the measurements were well below the detection limits for the diversion tank. Therefore, the amount of liquid lost cannot be determined accurately. Calculations made from field measurements and one sample analysis indicate that approximately 25 ft³ of soil were affected, and that 500 mCi of strontium-90, 500 mCi of cesium-137, and 130 mCi of cerium-144 were released with 12 gallons of liquid. The contaminated soil remains in place under 3 to 4 ft of clean soil.

31. Break in Underground Service Waste Line at ICPP

An underground service waste line carrying condensate from an evaporator failed at the point of transition from a 3-in. to a 2-in. line, at which there is a 90° change of direction. The break was discovered on September 30, 1976. Approximately 20,000 gallons of condensate leaked to the surrounding subsoil. The condensate contaminated approximately 38,700 ft³ of soil with 51 mCi of tritium, 2 mCi of strontium-90, 4 mCi of ruthenium-106, 2 mCi of cesium, and 1 mCi of cerium-144. The damaged section of line was excavated and repaired on October 2, 1976, and the contaminated soil covered with several feet of clean soil.

32. Pressurization of Solids Storage Cyclone at the ICPP Waste Calciner Facility

On October 26, 1976, the solids transport system at the Waste Calciner Facility (WCF) restricted, and analysis indicated that the solids storage cyclone was probably involved. During an attempt to clear the cyclone, it momentarily pressurized and blew out contaminated dolomite solids. The dolomite contaminated the roof of the vault and an area of approximately 300 ft² to the northeast of the waste calciner building. The release contained 230 mCi of strontium-90, 40 mCi of ruthenium-106, and 230 mCi of cesium-137. The majority of the contamination was removed from the vault rooftop and surrounding area. That remaining in the soil was covered with a layer of clean soil to fix it in place.

APPENDIX D
CALCULATIONAL METHODS

APPENDIX D

CALCULATIONAL METHODS

The calculational models and assumptions that were used in developing the radiological doses presented throughout the text of this statement are included in this appendix.

1. Whole Body Cloud Gamma Doses

The whole body cloud gamma doses were calculated by those methods described in Meteorology and Atomic Energy - 1968[D-1]. Radionuclides released from INEL facilities diffuse to large proportions, and by the time the material reaches offsite locations, the cloud (plume) can be considered infinite with respect to the distance gamma rays travel within the cloud. Based upon this consideration, it can be shown that the external whole body gamma dose to a receptor at ground level immersed in a plume of radionuclides is:

$$\text{Dose(rem)} = 0.25 \bar{E}_\gamma \Psi \quad (\text{D-1})$$

where

\bar{E}_γ = average gamma energy emitted at each disintegration
(MeV/dis.)

Ψ = concentration time integral of the cloud (Ci-sec/m³).

The concentration time integral was determined by multiplying the isotopic annual release by an annual atmospheric dispersion coefficient obtained by measured data at INEL. The atmospheric dispersion isopleths determined for 1974 are shown in Figure D-1. These isopleths are based on a continuous elevated release (250 ft) from the TRA-ICPP area.

The model (MESODIF) utilized to develop the isopleths is a refinement which avoids some of the previous deficiencies of the conventional wind rose technique [D-2]. The present model incorporates a boundary layer field of wind vectors in which spatial and temporal wind variabilities are included. Influences of local topography upon atmospheric dispersion are incorporated, somewhat indirectly but incorporated nonetheless, through simultaneous wind measurements at numerous locations over the Upper Snake River Plain.

The present model achieves two major improvements over the conventional wind rose technique. One improvement is the consideration of greater values of total integrated concentrations due to stagnation of airborne effluents over an area or systematic channeling and recurving of the windflow over the Upper Snake River Plain. The second major improvement results from an hour-by-hour adjustment of the rate of dilution when the hourly stability category changes.

NOTE: All values are $\times 10^{-8}$;
units are hr^2/m^3 .

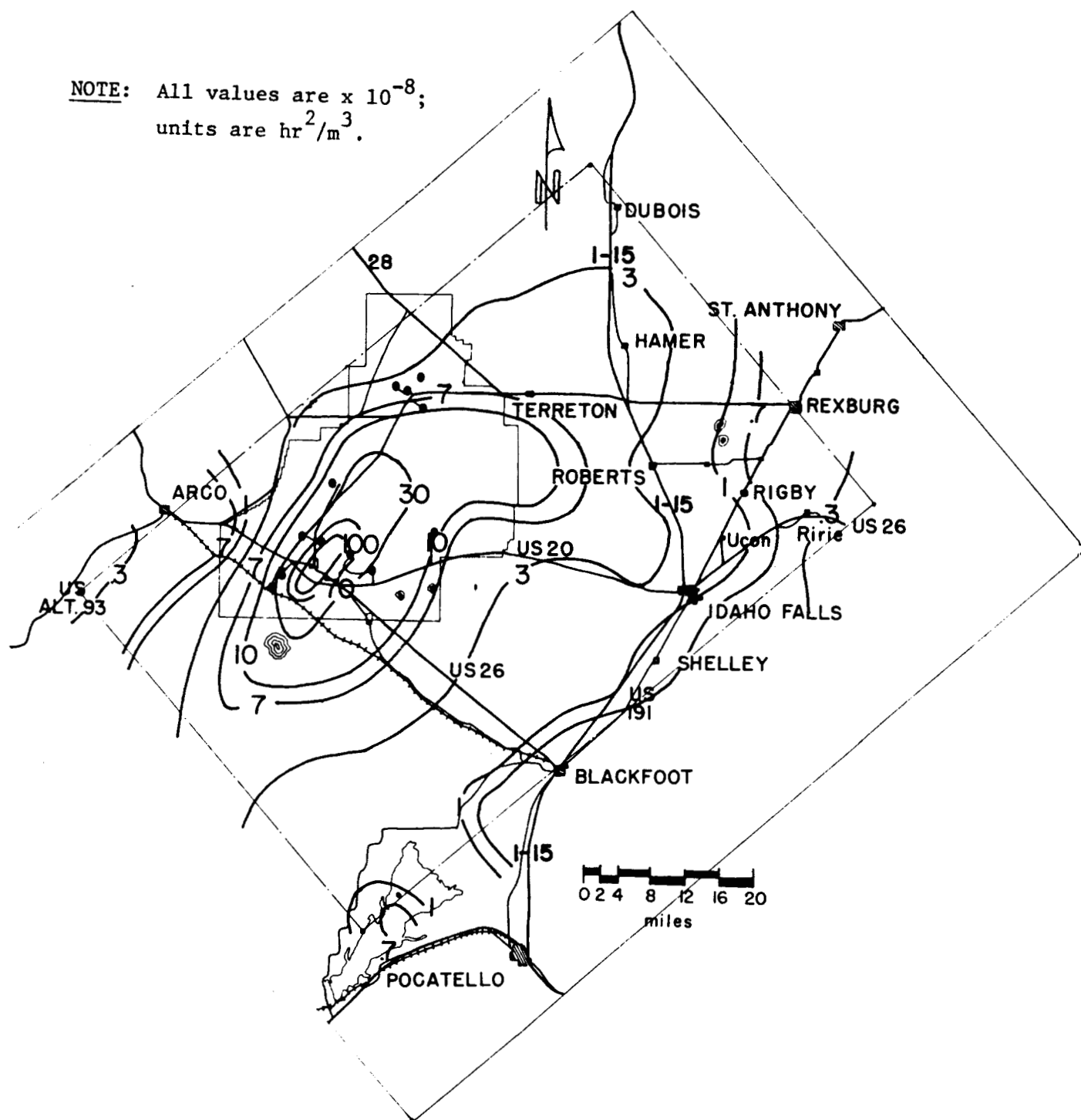


Figure D-1. Calculated Isopleths of Relative Annual Integrated Concentration for 1974.

Relative values of annual integrated concentrations for a postulated release from the TRA-ICPP complex during 1974 are shown in Figure D-1[a]. To obtain an annual time integrated air concentration (Ψ , Ci-hr/m³), the isopleth values (Ψ/Q) should be multiplied by the annual average hourly release rate (Q , Ci/hr).

The specific isopleth values for communities surrounding the INEL were used to calculate a 50-mile-radius population dose. The high isopleth value (Figure D-1) at the immediate southwest offsite location was used to calculate the maximum offsite dose.

2. Equation for Determining Radiation Dose from Inhalation or Ingestion of Radionuclides

The equation for determining the radiological doses from inhalation or ingestion of radionuclides was determined by the following standard equation:

$$\text{Radiation Dose (rad)} = \frac{P(3.7 \times 10^{10} \text{ d/s-Ci})(f)(E)(1.6 \times 10^{-6} \text{ ergs/MeV})(1-e^{-\lambda_e t})}{\lambda_e(m) (100 \text{ ergs/g-rad})} \quad (\text{D-2})$$

where

t = period of exposure (sec)

λ_e = biological decay constant (sec⁻¹)

P = uptake (Ci)

f = fraction of radionuclide reaching organ of reference

E = effective energy of radionuclide (MeV)

m = mass of organ of reference (g).

3. Method for Calculating Current and Cumulative Deposition on Soil

Figure D-1 was used to estimate the annual average air concentrations at locations on and near INEL. Using the average release rate (Q , Ci/hr), the annual time integrated air concentration (Ψ , Ci-hr/m³) was computed. The annual average air concentration (χ , Ci/m³) was also computed for use in computing the deposition rate (D , Ci/m²-day).

The deposition rate for worldwide fallout has been extensively studied in the United States. The following equation was developed to describe the deposition rate[D-3]:

[a] The concentrations were calculated assuming a release from the 250-ft-high (76-m) ICPP chimney. Plume rise was neglected in the calculations; therefore, the calculated ground level concentrations are likely to be slightly greater than what would have been measured.

$$D = (V_d + g \cdot r) \chi \quad (D-3)$$

where

- D = deposition rate ($\text{Ci}/\text{m}^2\text{-day}$)
 V_d = empirical constant for dry deposition of airborne particulates (m/day)
 g = empirical constant for precipitation scavenging of airborne particulates (m/cm of rain)
 r = rainfall rate (cm of rain/ day)
 χ = average air concentration (Ci/m^3).

The empirical values of V_d and g selected as appropriate for nuclear facility particulate releases were $V_d = 200 \text{ m}/\text{day}$ and $g = 4,530 \text{ m}/\text{cm}$ of rain[D-4]. The INEL average rainfall rate of $0.059 \text{ cm}/\text{day}$ was used in the calculations.

The equilibrium surface contamination S_e (Ci/m^2) was computed using

$$S_e = D/\lambda_r \quad (D-4)$$

where

λ_r is the radiological decay rate of the isotope (days^{-1}).

4. Method for Calculating Dose from Iodine-129 in Meat

The maximum annual average air concentration in areas open to grazing was obtained using Figure D-1 as described in Section 3 above. A release rate for iodine-129 of $0.1 \text{ Ci}/\text{yr}$ was assumed for each of the last 20 years. The median air-to-vegetation-surface-transfer velocity was taken to be $1 \text{ cm}/\text{sec}$ or $864 \text{ m}/\text{day}$ for dry processes[D-5] and an upper limit value of $14,000 \text{ m}/\text{cm}$ of rain[D-3] was assumed for wet processes. The deposition rate onto vegetation is then

$$D_v = (864 + 14,000 r) \chi \quad (\text{Ci}/\text{m}^2\text{-day}) \quad (D-5)$$

and the equilibrium concentration on vegetation (S_v , Ci/m^2) will be

$$S_{v_e} = D_v/\chi_e$$

where χ_e is the effective removal rate (days^{-1}) for radiiodine on vegetation. A value of λ_e of 0.05 days^{-1} was assumed.

It is estimated that the transfer parameter for ingested iodine-129 to meat is $2\%/ \text{kg}$ [D-6]. It was assumed that grazing animals consumed a maximum of $12 \text{ kg}/\text{day}$ (dry weight) and that the forage density was $0.16 \text{ kg}/\text{m}^2$ (dry weight). The effective half-life of iodine-129 in meat was taken to be 33 days so, after a grazing period of 90 days, the concentration in meat would be $\sim 90\%$ of the equilibrium value.

An adult is assumed to consume 80 kg of meat per year, and the dose conversion factor of 7.05×10^{-3} mrem/pCi ingested was utilized. An additional 1% of the dose per year of accumulation to account for soil-root transfer was assumed[D-6].

5. Method for Calculating External Dose from Equilibrium Deposition of Gamma Emitters in Soil

The equilibrium deposition was calculated as described in Section 3 above. The exposure rate at 1 m in air was computed using data in Table 9 of Reference D-7. The relaxation length α was assumed to be 2 cm; the soil density ρ was assumed to be 1.6 g/cm³. A quality factor of 1 and the rad/R conversion factors from Reference D-8 were used. To determine the genetically significant dose, the multiplicative body screening and dwelling shielding factors of 0.8 and 0.4[D-4] were employed.

6. Method for Calculating Doses from Iodine-131 and Iodine-129 in Milk

Reference D-9 states that a continuous air concentration of 4.2×10^{-15} $\mu\text{Ci}/\text{cm}^3$ of elemental iodine-131 gas will produce a milk concentration of 2.4 pCi/liter and a resulting dose of 15 mrem/yr to the thyroid of an infant drinking 1 liter of milk per day. Potential doses to 1-year-old children around INEL were computed using that concentration/dose relationship. No credit for decay in processing or transit was assumed; that is, the milk was assumed to be taken directly from a family cow. Dose conversion factors for iodine-129 ingestion were taken from Reference D-6. In both cases, the possible dose from milk consumption was based on the 1974 milk monitoring data.

7. Method for Calculating Doses from Ingestion of Cesium-137 in Meat

The deposition of cesium-137 from worldwide fallout has been correlated with the concentration of cesium-137 in milk and the data fit the equation:

$$C_m = 3.6 [D(n) + 0.65 D(n-1) + D(n-2)] \quad (\text{D-7})$$

where C_m is the average milk concentration in pCi/liter in year n , and $D(n)$, $D(n-1)$, and $D(n-2)$ are the annual deposition rates $\mu\text{Ci}/\text{m}^2$ in years n , $n-1$, and $n-2$, respectively[D-10].

It was also observed that the ability of cesium-137 in meat was consistently related to that in milk and that the following ratio held.

$$\frac{(\text{pCi/kg of meat})}{(\text{pCi/liter of milk})} \sim 4. \quad (\text{D-11}) \quad (\text{D-8})$$

The maximum equilibrium deposition rate for cesium-137 released from INEL facilities was calculated as described in Section D.3 to be $1.43 \mu\text{Ci}/\text{m}^2\text{-yr}$; this value was used for $D(n)$, $D(n-1)$, and $D(n-2)$ in the computation of the equilibrium meat concentration.

Maximum equilibrium body burdens and doses for four age groups were computed using the meat consumption rates given in Reference D-6 and the age dependent dose conversion factors developed in Reference D-4. The highest postulated dose was received by an adult. For those cases when doses were computed based on assumed consumption of meat with known cesium-137 concentrations, the same dosimetric parameters were used with the known concentration.

8. References

- D-1. D. H. Slade (ed.), Meteorology and Atomic Energy - 1968, TID-24190 (July 1968).
- D-2. G. E. Start and L. L. Wendell, Regional Effluent Dispersion Calculations Considering Spatial and Temporal Meteorological Variations, NOAA Technical Memorandum ERL ARL-44 (May 1974).
- D-3. C. A. Pelletier and P. G. Voilleque, "The Behavior of Cesium-137 and Other Fallout Radionuclides on a Michigan Dairy Farm," Health Physics, 21, 777 (1971).
- D-4. P. G. Voilleque and C. A. Pelletier, "Comparison of External Irradiation and Consumption of Cows' Milk as Critical Pathways for Cs-137, Mn-54, and Ce-Pr-144 Released to the Atmosphere," Health Physics 27, 189 (1974).
- D-5. J. D. Zimbrick and P. G. Voilleque (eds.), Controlled Environmental Radioiodine Tests at the National Reactor Testing Station, Progress Report Number Four, USAEC Report IDO-12065 (January 1969).
- D-6. J. K. Soldat, Environmental Behavior and Radiation Doses from I-129, USAEC Report BNWL-SA-4879 (June 1974).
- D-7. H. L. Beck et al, In Situ Ge(Li) and NaI(Tl) Gamma Ray Spectrometry, USAEC Report HASL-258 (September 1972).
- D-8. The International Commission on Radiation Units and Measurements, Physical Aspects of Irradiation, NBS Handbook 85, Washington, D. C.: U. S. Government Printing Office (1964).
- D-9. USAEC Directorate of Regulatory Standards, Regulatory Guide 1.42, Interim Licensing Policy on as Low as Practicable for Gaseous Radioiodine Releases from Light-Water-Cooled Nuclear Power Reactors (June 1973).
- D-10. Report of the United Nations Scientific Committee on the Effects of Atomic Radiation, U. N. General Assembly Official Records: Nineteenth Session, Supplement No. 14 (A/5814), New York: United Nations (1964).

D-11. Report of the United Nations Scientific Committee on the Effects of Atomic Radiation, U. N. General Assembly Official Records: Twenty-First Session, Supplement No. 14 (A/6314), New York: United Nations (1966).